

Lee Boyling, BSc (Hons)

Head of *In Vitro/Ex vivo* Study Direction

Alnwick, UK

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Key Roles and Responsibilities

- ▶ Management of team conducting drug metabolism, exploratory bioanalysis and *in vitro* biochemical assays
- ▶ Member of Global Metabolism Team
- ▶ Member of Alnwick Safety Assessment *In Vivo* Study Director team

Areas of Expertise

- ▶ **Drug Metabolism**
 - Biotransformation
 - Design, conduct and interpretation of metabolism studies
 - Support of investigative toxicology studies
- ▶ **Bioanalysis**
 - Conduct of exploratory LC-MS bioanalysis
 - Study Director for GLP Bioanalysis
- ▶ **Drug Development**
 - Wide experience from entry into development up to Human AME

Professional Highlights

- ▶ More than 20 years experience within the pharmaceutical industry
- ▶ Joined Sterling Winthrop (latterly sanofi-aventis) in 1992 as a scientist within Drug Metabolism, Study Directing *in vitro* CYP activity assays

- ▶ By 1996 progressed to study directing nonclinical *in vivo* mass balance studies
- ▶ In 2000 changed to Clinical Metabolism which included working alongside Clinical Pharmacology to organize human AME studies
- ▶ In 2004 was lead scientist responsible for development of LC-MS metabolomics platform and in 2006 was Company Representative on COMET2 consortium
- ▶ Following acquisition of Alnwick site by Covance in November 2010, became Head of Metabolism at Alnwick in June 2011
- ▶ In November 2014 became Head of *In Vitro/Ex vivo* Study Direction formed by merging Alnwick Drug Metabolism and Molecular Cellular Toxicology groups
- ▶ Lecturer on external course (UK Drug Metabolism Discussion Group Fundamentals Training Course)

Education

- ▶ **BSc, Applied Biology**
University of Hertfordshire

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